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## Multistate Outbreak of Listeriosis — United States, 2000

Since May 2000, 29 illnesses caused by a strain of *Listeria monocytogenes* (LM) have been identified in 10 states: New York (15 cases); Georgia (three); Connecticut, Ohio, and Michigan (two each); and California, Pennsylvania, Tennessee, Utah, and Wisconsin (one each). Dates of LM isolation ranged from May 17 through November 26 with 26 (90%) infections occurring since July 15. When subtyped, the LM isolates from these cases were indistinguishable by pulsed-field gel electrophoresis (PulseNet pattern numbers GX6A16.0014 by *Asc*1 and GX6A12.0017 by *Apa*1) and ribotyping (DUP-1053). This report summarizes the investigation, which linked these cases of listeriosis to eating deli turkey meat.

Eight perinatal and 21 nonperinatal cases were reported. Among the 21 nonperinatal case-patients, the median age was 65 years (range: 29–92 years); 13 (62%) were female. The 29 cases have been associated with four deaths and three miscarriages/stillbirths.

A case-control study conducted by five state and two local health departments and CDC implicated eating deli turkey meat as the probable source of infection. Thirteen (76%) of 17 case-patients and five (21%) of 24 controls ate deli turkey meat during the 30 days before illness onset (Mantel-Haenszel weighted odds ratio=8.0; 95% confidence interval=1.2–43.3). State health and agriculture departments investigated 13 stores and delicatessens where 11 patients reported purchasing turkey; these stores and delicatessens carried turkey meat produced by at least 27 federally inspected establishments. Two establishments were linked to 10 of 11 patients; one of these establishments produced turkey meat for the second establishment.

On December 8, investigators from the Food Safety and Inspection Service, U.S. Department of Agriculture (USDA) began investigating the implicated establishments. On December 12, Cargill Turkey Products, Inc. (Waco, Texas) stopped shipping ready-to-eat foods and, on December 14, voluntarily recalled processed turkey and chicken deliment that might have been contaminated.

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Editorial Note: LM infection causes an estimated 2500 serious illnesses and 500 deaths in the United States each year. Infected pregnant women may experience only a mild, influenza-like illness; however, infections during pregnancy can lead to premature delivery, miscarriage, stillbirth, or serious infection of the newborn. Other persons at increased risk for infection are those aged ≥65 years, persons with cancer, diabetes, kidney disease, acquired immunodeficiency syndrome, or who take immunosuppressive medications. Manifestations of illness include meningitis and sepsis. Healthy persons aged <65 years rarely are affected.

The risk for a person developing *Listeria* infection after eating a contaminated product is very small. Persons who have eaten a recalled product but do not have symptoms do not require tests or treatment even if they are in a high-risk group. However, persons in a high-risk group who have eaten contaminated product and become ill within 2 months with fever or signs of serious illness should consult a physician.

Guidelines for preventing listeriosis are similar to those for preventing other foodborne illnesses. The general recommendations are 1) cook thoroughly raw food from animal sources (e.g., beef, pork, or poultry); 2) wash raw vegetables thoroughly before eating; 3) keep uncooked meats separate from vegetables and from cooked foods and ready-to-eat foods; 4) avoid raw (unpasteurized) milk or foods made from raw milk; and 5) wash hands, knives, and cutting boards after each handling of uncooked foods. Persons at high risk for listeriosis may choose to 1) avoid soft cheeses (i.e., feta, Brie, Camembert, blue-veined, and Mexican-style cheese such as queso fresco). Hard cheeses, processed cheeses, cream cheese, cottage cheese, or yogurt need not be avoided; 2) cook leftover foods or ready-to-eat foods (e.g., hot dogs) until steaming hot; and 3) avoid foods from deli counters (e.g., prepared salads, meats, and cheeses) or thoroughly reheat cold cuts before eating.

Cases of listeriosis with onset since October 1, 2000, should be reported to state and local health departments; information about the recall is available at http://www.fsis.usda.gov/OA/recalls/rec\_actv.htm\*. Consumers who have recalled meat products, even if they have been stored in freezers, should discard or return them to the point of purchase. High-risk consumers who have processed turkey or chicken deli meat but are uncertain of the brand should call the place of purchase to find out if it might be a recalled product, or discard it. Answers to meat-safety questions are available at the USDA meat and poultry hotline, (800) 535-4555. Listeriosis information is available at http://www.cdc.gov/ncidod/dbmd/diseaseinfo/listeriosis\_g.htm.

<sup>\*</sup>References to sites of non-CDC organizations on the World-Wide Web are provided as a service to MMWR readers and do not constitute or imply endorsement of these organizations or their programs by CDC or the U.S. Department of Health and Human Services. CDC is not responsible for the content of pages found at these sites.

## Foodborne Outbreak of Group A Rotavirus Gastroenteritis Among College Students — District of Columbia, March-April 2000

1131

On March 31, student health services at a university in the District of Columbia (DC) notified the DC health department that an increased number of students had become ill with acute gastroenteritis beginning March 29. Some ill students reported eating tuna or chicken salad sandwiches from dining hall A on campus. On March 31, the DC health department initiated an outbreak investigation. This report summarizes results of the investigation, which indicated that group A rotavirus transmitted by food was the cause of the outbreak.

Telephone interviews were conducted with students who reported illness to student health services, with additional ill students who were identified during interviews, and with healthy controls selected randomly from the university registry of students residing on campus. A case of gastroenteritis was defined as three or more episodes of diarrhea and/or two or more episodes of vomiting within a 24-hour period in a student with onset on or after March 20. Controls and case-patients whose illness onset occurred during March 27–31 were questioned about food history, residence and dining hall, source of water, use of a public access computer or sports equipment at the university gym, and attendance at social or athletic events. Electronic records of student meal attendance were available for 49 case-patients with illness onset during March 27–31 and for 55 control subjects.

Twenty-three (79%) of 29 employees of dining hall A were interviewed to identify their work duties and determine whether they were ill. Stool specimens were collected during March 29–April 10 from six ill students and 21 dining hall A employees. Samples were screened for bacterial and parasitic pathogens at a commercial laboratory and for viral pathogens at CDC.

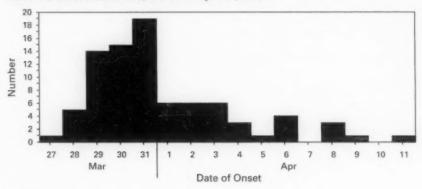
The outbreak among students began March 27 and peaked at 19 cases on March 31 (Figure 1). A total of 108 students (55 were identified by telephone interviews and 53 were self-reported) had gastrointestinal symptoms during March 26–April 11; 85 (79%) had illness that met the case definition. The attack rate among students residing on campus was 5% (77 of 1641), with no significant differences in attack rates by sex, occupancy of residence hall, or grade level. Eight case-patients resided off campus (attack rate: 0.02%). Among the 83 case-patients for whom a complete list of symptoms was reported, 77 (93%) had diarrhea, 75 (90%) abdominal pain or discomfort, 69 (83%) loss of appetite, 67 (81%) nausea, 64 (77%) fatigue, 56 (67%) vomiting, 49 (59%) headache, 48 (58%) chills, 48 (58%) subjective or low-grade fever, and 42 (51%) myalgia. Sore throat, cough, and/or congestion were reported by six case-patients with onsets on or after April 2. The median duration of illness was 4 days (range: 1–8 days). Nine (11%) case-patients received intravenous fluids to treat dehydration.

Of those who completed the telephone interview, 40 (91%) of 44 case-patients and 27 (68%) of 40 controls ate at least one deli sandwich from campus dining hall A during March 27–30 (p=0.017; odds ratio [OR]=4.8; 95% confidence interval [CI]=1.3–22.1). During March 27–30, four (8%) of 49 case-patients ate four or more meals at dining hall B compared with 18 (33%) of 55 controls (p=0.005; OR=0.2; 95% CI=0.04–0.6). Food histories of employees were not recorded; however, six employees reported illness.

Stool specimens of students and employees were negative for bacterial and parasitic pathogens and for Norwalk-like viruses. Using electron microscopy, enzyme immunoassay, and reverse transcriptase-polymerase chain reaction (RT-PCR), nine (33%) of 27

Rotavirus Gastroenteritis - Continued

FIGURE 1. Number\* of gastroenteritis\* cases among college students, by date of illness onset — District of Columbia, March 27–April 11, 2000



# n=85

A case of gastroenteritis was defined having three or more episodes of diarrhea and/or two or more episodes of vomiting within a 24-hour period in a student with onset on or after March 20.

specimens were positive for group A rotavirus. Rotavirus positive stool specimens from four students and three employees were identified as genotype combination P[4],G2 by RT-PCR. Two of the three P[4],G2-positive employees were line cooks who reported having symptoms of gastroenteritis on March 27 and April 2, respectively, while the third positive employee, a deli server, reported no illness.

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Editorial Note: Group A rotavirus is the most common cause of childhood diarrhea worldwide, infecting >90% of children by age 3 years (1). Because rotavirus immunity develops early in life, disease among older children and adults is uncommon (1). Although the role of rotavirus in diarrhea outbreaks in adults has not been well studied, it has been documented as the cause of adult diarrheal outbreaks in hospitals (2), nursing homes (3), isolated communities (4), and in travelers (5). Also, parents of children infected with rotavirus have been reported to experience acute gastroenteritis (6). However, the rotavirus G and P protein-type combinations, the proteins that elicit an immune response in humans, were not characterized in most of these reports.

The rapid increase and gradual decline of the campus outbreak suggest that the infection was foodborne during the first week and was spread person-to-person during the following week. During the first week, illness was associated with eating sandwiches at dining hall A and was associated inversely with eating frequently at dining hall B. The employee who prepared sandwich fillings did not report illness and tested negative for rotavirus. None of the three deli servers who assembled and served sandwiches reported illness; however, one was rotavirus P[4],G2 positive. It is unknown whether the deli server who tested positive was infected before the outbreak among students.

#### Rotavirus Gastroenteritis — Continued

This rotavirus serotype G2 outbreak was unusual for two reasons; food was implicated as the source of infection and the adults affected should have been immune. During April 2000, a gastroenteritis outbreak among adults in Japan also was caused by foodborne transmission of group A rotavirus serotype G2 (7). These adults should not have been susceptible to severe rotavirus illness. G2 strains often are found combined with serotype P[4]1B (8). The G and P neutralization antigens of serotype G2 strains may allow G2 strains to escape immunity induced by the more common G1, G3, and G4 strains. In addition, G2 has been associated with more severe dehydration during diarrheal episodes in children than other common strains (9). These outbreaks of rotavirus gastroenteritis in adults in the United States and Japan raise questions about the persistence of immunity to rotavirus and the virulence of G2 strains. Investigators and clinicians should consider rotavirus as a possible cause of acute gastroenteritis in adults.

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## Blood Lead Levels in Young Children — United States and Selected States, 1996–1999

Lead exposure adversely affects the cognitive development and behavior of young children (1). For children aged <6 years, CDC has defined an elevated blood lead level (BLL) as ≥10 µg/dL, but evidence exists for subtle effects at lower levels (2). Data from CDC's Third National Health and Nutrition Examination Survey, Phase 2 (1991–1994) (NHANES) showed that average BLLs in children had decreased approximately 80% since the late 1970s but that elevated BLLs remained more common among low-income children, urban children, and those living in older housing (3,4). Although these data provide national estimates of the prevalence of elevated BLLs among children, they do not provide information at the state or local level. To target prevention efforts and monitor progress toward reducing BLLs at the state and local level, CDC's Childhood Blood Lead Surveillance (CBLS) program supports state blood lead surveillance programs on the basis of blood lead tests from public and private clinical laboratories. This report

summarizes data on BLLs in children aged 1–5 years from NHANES data collected in 1999 and children aged <6 years from state surveillance data provided to CDC by 19 state surveillance programs during 1996–1998. The findings indicate that, despite the decreases in mean BLL among children, the problem remains concentrated on a local level. Surveillance efforts should be used to target screening efforts to communities at highest risk.

NHANES is a continuous survey of the health and nutritional status of the U.S. civilian, noninstitutionalized population designed so that each year of data constitutes a nationally representative sample. Data in this report are from NHANES 1999, and NHANES III, Phase 2. A household interview and a physical examination were conducted for each survey participant. During the physical examination, blood was collected by venipuncture for all persons aged >1 year. Graphite furnace atomic absorption spectrophotometry was used to measure BLLs with detection limits of 0.3 µg/dL (NHANES 1999) and 1.0 µg/dL (NHANES III, Phase 2). Long-term quality-control data for these analyses, including similar standardized reference materials, were used in both surveys and showed that data from the two surveys can be compared. Because of limited sample size, NHANES 1999 analyses include only data on average BLLs and selected percentiles but not on the prevalence of elevated levels.

The analyses of CBLS data were based on reports from 19 of 28 states that provided blood lead data to CDC (Table 1). The 19 states were included because they received all blood lead test results of children from participating laboratories (regardless of level) and reported data from January 1, 1996 through December 31, 1998. These states accounted for 33% of all U.S. children aged <6 years.

An elevated BLL from CBLS is defined as a single blood lead test result  $\geq 10~\mu g/dL$ . If multiple tests were reported for a child during a calendar year, the highest BLL measured for that child was used. To estimate the proportion of children with elevated BLLs among those tested, the number of children with elevated levels was divided by the number of children tested at least once during a calendar year.

From NHANES III, Phase 2 (1991–1994) to NHANES 1999, the geometric mean BLL in children aged 1–5 years decreased from 2.7 (95% confidence interval [CI]=2.6–2.9) to 2.0  $\mu$ g/dL (95% CI=1.7–2.3), and the 50th percentile decreased from 2.6 (95% CI=2.4–2.8) to 1.9  $\mu$ g/dL (95% CI=1.6–2.1). The continued pattern of decline in BLLs between the two surveys also is indicated at the 10th, 25th, 75th, and 90th percentiles.

The CBLS data showed that the proportion of children tested with BLLs  $\geq$ 10 µg/dL decreased from 10.5% in 1996 to 7.6% in 1998 in the 19 states providing data (Table 1). The proportions of children with BLLs  $\geq$ 15 µg/dL and  $\geq$ 20 µg/dL also decreased.

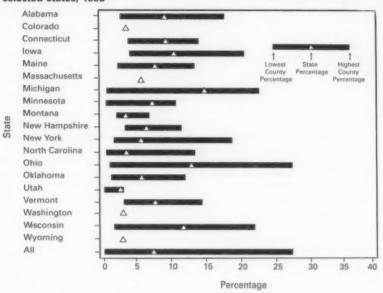
The percentage of children aged <6 years tested with BLLs ≥10 µg/dL in each state ranged from 2.7 to 14.9 (Figure 1). Within states, the proportion of children with elevated

TABLE 1. Percentage of children tested aged <6 years with elevated blood lead levels (BLLs), by year — selected states\*, 1996–1998

		% Children	n with elevated BL	Ls (µg/dL)
Year	No. tested	≥10	≥15	≥20
1996	1,220,596	10.5%	3.9%	1.9%
1997	1,183,506	8.6%	3.2%	1.5%
1998	1,256,907	7.6%	2.7%	1.2%

<sup>\*</sup> Alabama, Colorado, Connecticut, Iowa, Maine, Massachusetts, Michigan, Minnesota, Montana, New Hampshire, New York, North Carolina, Ohio, Oklahoma, Utah, Vermont, Washington, Wisconsin, and Wyoming.

FIGURE 1. State-specific percentage of children aged <6 years tested with blood lead levels (BLLs)  $\geq$ 10  $\mu$ g/dL and highest and lowest percentage of elevated BLLs, by county — selected states, 1998\*



\* Only counties with ≥200 children tested for BLL are included. Colorado, Washington, and Wyoming had <2 counties with 200 children tested, and Massachusetts did not report county of residence.

BLLs in counties with at least 200 children tested also varied considerably. For example, the proportion of children with elevated BLLs ranged from 1.3% to 27.3% in counties in Ohio. Across all 19 states, the county-specific proportions of children with elevated BLLs ranged from 0.5% to 27.3%, indicating a concentrated proportion of elevated BLLs in specific populations or geographic areas.

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Editorial Note: The findings in this report indicate that average BLLs of U.S. children aged 1-5 years have declined from the early 1990s to 1999. Because of the limited sample size of a single year of NHANES 1999 compared with that of the multiple years of NHANES III, additional data are necessary to confirm this trend. The dramatic decline in BLLs from the late 1970s through the early 1990s resulted primarily from the phaseout of leaded gasoline and the resulting decrease in lead emissions, although other exposures also decreased (3). Although air lead levels and lead emissions continued to decrease during the 1990s, most of this decline occurred before 1995 (5). The primary remaining sources of childhood lead exposure are deteriorated leaded paint and the soil and dust it contaminates in old housing. The construction of new housing and the demolition and rehabilitation of older housing may be contributing to a continued decline in BLLs. Data from NHANES III, Phase 2 showed that low-income children living in older housing had more than a 30-fold greater prevalence of BLLs ≥10 µg/dL than do middleincome children in newer housing (4). From 1993 to 1997, the number of low-income children living in pre-1940s and 1940-1974 housing declined by 31% and 14%, respectively. The number of low-income children living in post-1974 housing increased by 5% (6).

Despite the overall decline in average BLLs, CBLS data show that the risk for elevated BLLs in children tested remains high in some counties and varies greatly among and within states. This variation most likely reflects geographic variation in the prevalence of risk factors for elevated BLLs such as residence in older housing and poverty.

The findings in this report are subject to at least four limitations. First, the small NHANES 1999 sample does not permit observing risks in specific subgroups or geographic areas, but it provides a nationally representative estimate of BLLs in children. The CBLS data set provides local information but is limited to children who receive clinical or diagnostic blood lead testing. Second, because CDC guidelines recommend the use of blood lead data and census data to target screening efforts in populations at increased risk for lead exposure, the proportion of children with elevated BLLs is higher in CBLS data than would be expected in NHANES 1999. Third, the guidelines for testing children vary by state, and adherence to the guidelines varies by health-care provider. Finally, CBLS data include samples collected by fingerstick, which can slightly overestimate the blood lead result, and venous samples and results obtained by different laboratories. Despite these differences, the temporal trends in BLLs are similar between the CBLS and NHANES data sets.

One of the national health objectives for 2010 is the elimination of childhood lead poisoning (7). Data in this report document continued progress toward this goal but also show the ongoing need to target prevention efforts at communities and populations at highest risk. CDC recommends that state health agencies target screening efforts by using blood lead surveillance data, census data, Medicaid data, and other sources of information on risk factors such as housing age and poverty (8,9). Other federal agencies, including the U.S. Department of Housing and Urban Development and the U.S.

Environmental Protection Agency, also are implementing targeted strategies to prevent lead exposure. State blood lead surveillance systems play a key role in targeting and monitoring federal, state, and local prevention efforts. CDC encourages additional states to conduct surveillance for elevated BLLs in children.

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## Notice to Readers

# Public Health Service Recommendations for the Use of Vaccines Manufactured with Bovine-Derived Materials

The Center for Biologics Evaluation and Research (CBER), U.S. Food and Drug Administration (FDA) learned earlier this year that some vaccines were manufactured with bovine-derived materials obtained from countries in which bovine spongiform encephalopathy (BSE) or a substantial risk for BSE exists. A list of these countries is published by the U.S. Department of Agriculture (USDA).\* This information was of concern because cases of variant Creutzfeldt-Jakob disease (vCJD) have been attributed to, among other possibilities, eating beef products from cattle infected with the agent of BSE. No evidence exists that cases of vCJD are related to the use of vaccines, and no cases of vCJD have been reported in the United States.

CBER assessed the risk for vCJD from vaccines manufactured with processes that use bovine materials potentially contaminated with the BSE agent. On July 27, 2000, CBER convened a joint meeting of the Transmissible Spongiform Encephalopathy Advisory Committee and the Vaccines and Related Biological Products Advisory Committee to review the results of these assessments and make recommendations about the use and manufacture of these vaccines. The committees concluded that the risk for vCJD

<sup>\*9</sup> CFR, part 94.

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posed by vaccines in the scenarios presented was theoretical and remote. This conclusion was based on the inherent low risk of the bovine materials involved (e.g., type and amount of tissue[s] used, specific time and country, or herd of origin) and/or the dilutions of materials during manufacture. The committees concluded that the benefits of vaccination outweigh any remote risks for vCJD.

As a precautionary measure, the committees recommended that vaccines manufactured with bovine-derived materials from countries on the USDA list be replaced with bovine-derived materials from other countries. This recommendation, which is consistent with existing FDA guidance first issued in 1993 on the sourcing of bovine-derived materials, is intended to reduce even the remote risk for vCJD from vaccines. The committees also recommended that FDA provide information to the public about the safety of vaccines made with materials from countries in which BSE or BSE risk exists.

FDA has requested that manufacturers replace bovine-derived materials obtained from countries on the USDA list with materials obtained from countries not on the USDA list. All of the affected manufacturers have agreed to implement these changes or have already done so. FDA anticipates that most of these changes will be completed in 2001.

The Public Health Service (PHS) recommends that all persons continue to be vaccinated according to current schedules. PHS has no preference for using one licensed vaccine product over another based on the source of bovine-derived materials used in vaccine production. Failure to obtain the recommended vaccinations with licensed vaccines poses a risk for serious disease.

Additional information about BSE or vaccines manufactured with bovine-derived materials from countries on the USDA list can be obtained from the FDA World-Wide Web site, http://www.fda.gov/cber/BSE/BSE.htm¹, or from CBER's Office of Communications, Training and Manufacturers Assistance, telephone (800) 835-4709.

## Notice to Readers

# Availability and Use of Parenteral Quinidine Gluconate for Severe or Complicated Malaria

Since 1991, quinidine gluconate, a class 1a anti-arrhythmic agent, has been the only parenteral antimalarial available for use in the United States (1). It is indicated for the treatment of patients with life-threatening *Plasmodium falciparum* malaria (2), including those who cannot tolerate oral therapy, have high-grade parasitemia, or have complications (e.g., cerebral malaria or acute renal failure) (3,4).

The limited availability of and delays in obtaining quinidine gluconate have contributed to adverse patient outcomes (5–7). As newer anti-arrhythmics have replaced quinidine for many cardiac indications, some hospitals and other health-care facilities have dropped quinidine gluconate from their formularies and, as a result, fewer clinicians have had experience using the drug. Discussions among quinidine gluconate manufacturer Eli Lilly Company (Indianapolis, Indiana), CDC, the U.S. Department of Defense, and the U.S. Food and Drug Administration have resulted in the following recommendations

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to improve quinidine gluconate availability for acutely ill malaria patients in U.S. healthcare facilities:

- Before an acute need arises, hospital drug services should consider maintaining or adding quinidine gluconate to formularies or should be able to immediately locate a nearby source.
- Pharmacists and clinicians requiring quinidine gluconate in hospitals in which an immediate source cannot be located should contact their local or regional distributor to request quinidine gluconate.
- 3. In clinical settings in which the need for the drug is more acute than can be met by the local or regional distributor, pharmacists and clinicians should contact Eli Lilly Company, telephone, (800) 821-0538 to arrange a rapid shipment of the drug. This telephone number, or an alternate number given to callers, is staffed 24 hours a day, 7 days a week.
- 4. If further assistance is needed in obtaining quinidine gluconate or in managing patients with malaria, contact CDC's malaria hotline, (770) 488-7788 (Monday–Friday, 8 a.m. to 4:30 p.m. eastern standard time). After business hours, weekends, and holidays, contact CDC's security station, telephone, (404) 639-2888 and ask to have the on-call person for malaria questions paged.

The following dosing recommendations for quinidine gluconate administration are provided for pharmacists and clinicians treating patients with severe or complicated malaria:

- Quinidine gluconate intravenous should be administered in a monitored setting.
   Prolongation of the QT interval as indicated by an electrocardiogram, ventricular arrhythmia, hypotension, and hypoglycemia can result from the use of this drug at treatment doses.
- Quinidine gluconate for malaria is administered as an initial intravenous loading dose of 10 mg/kg salt (equivalent to 6.25 mg/kg quinidine base) infused over 1–2 hours. Quinidine gluconate is administered subsequently as a continuous infusion of 20 µg/kg/min quinidine gluconate salt (equivalent to 12.5 µg/kg/min quinidine base) (2).
- An alternative regimen is an intravenous loading dose of 24 mg/kg quinidine salt (equivalent to 15 mg/kg quinidine base) infused over 4 hours, followed by a maintenance infusion of 12 mg/kg of quinidine gluconate salt (equivalent to 7.5 mg/kg quinidine base) infused over 4 hours every 8 hours, starting 8 hours after the loading dose (2). These regimens have been shown to be effective with or without concomitant exchange transfusion (2).
- The risk for serious ventricular arrhythmia associated with quinidine is increased by bradycardia, hypokalemia, and hypomagnesemia (2). When determining whether a patient should receive a bolus dose, previous administration of other drugs that can prolong the QT interval (e.g., quinine, halofantrine, and mefloquine) should be considered.
- No alternatives to quinidine exist for patients in the United States who require intravenous therapy for malaria. Acute cardiac events can be minimized by careful calculation of the loading dose and infusion rate. Consulting a cardiologist may be helpful when attempting to resume infusion in the patient who has experienced QT prolongation or hypotension associated with intravenous quinidine infusion.
- · Consulting a physician with experience in treating malaria is advised.

#### Notices to Readers - Continued

#### References

- CDC. Treatment with quinidine gluconate of persons with severe Plasmodium falciparum infection: discontinuation of parenteral quinine from CDC drug service. MMWR 1991;40(no. RR-4):21–3.
- Quinidine gluconate injection [package insert]. Indianapolis, Indiana: Eli Lilly Company, February 2000.
- Zucker JR, Campbell CC. Malaria: principles of prevention and treatment. Infect Dis Clin No. Am 1993:7:547–67.
- Miller KD, Greenberg AE, Campbell CC. Treatment of severe malaria in the United States with a continuous infusion of quinidine gluconate and exchange transfusion. N Engl J Med 1989;321:65–70.
- Rosenthal PJ, Petersen C, Geertsma FR, et al. Availability of intravenous quinidine for falciparum malaria (Letter). N Engl J Med 1996;335:138.
- Humar A, Sharma S, Zoutman D, et al. Fatal falciparum malaria in Canadian travelers. Can Med Assoc J 1997:156:1165–7.
- CDC. Availability of parenteral quinidine gluconate for treatment of severe or complicated malaria. MMWR 1996;45:494–5.

### Notice to Readers

## Availability of MMWR Mirror Website in Spain

CDC, in collaboration with the Toxic Oil Syndrome Research Centre (CISAT) of the Institute of Health Carlos III, Madrid, Spain, has established a MMWR mirror website in Spain. The website was developed to reduce the delay caused by transoceanic electronic transfers of large documents and to increase access to information published in MMWR for European public health practitioners. The mirror website is updated simultaneously with the posting of new reports on the MMWR website (http://www.cdc.gov/mmwr). The address for the CISAT MMWR mirror website is http://cisat.isciii.es/mmwr. The website also hosts a mirror site from the Agency for Toxic Substances and Disease Registry (ATSDR). This mirror site can be found at http://cisat1.isciii.es/atsdr. Other features of the website include information on environmental health problems and rare diseases in Spanish.

CISAT is a part of the WHO Collaborating Centre for the Clinical Epidemiology of Environmental Diseases and has established agreements with CDC/ATSDR and the University of Pittsburgh. Support of the *MMWR* mirror website is part of a larger effort undertaken by CISAT to create a comprehensive environmental health information site.

# Notice to Readers

# **Epidemiology in Action: Intermediate Methods**

CDC and Emory University's Rollins School of Public Health will co-sponsor a course, "Epidemiology in Action: Intermediate Methods" during February 26–March 2, 2001, at Emory University. The course is designed for state and local public health professionals.

#### Notices to Readers - Continued

The course will review the fundamentals of descriptive epidemiology and biostatistics, analytic epidemiology and computers as used in epidemiology but will focus on midlevel epidemiologic methods directed at strengthening participants' quantitative skills, with an emphasis on up-to-date data analysis. Topics include field investigations, advanced measures of association, normal and binomial distributions, logistic regression, and additional statistical methods. Prerequisite is an introductory course in epidemiology, such as Epidemiology in Action, International Course in Applied Epidemiology or any other introductory class. There is a tuition charge.

Deadline for applications is January 15. Additional information and applications are available from Emory University, Rollins School of Public Health, International Health Dept. (PIA), 1518 Clifton Road, N.E., Room 746, Atlanta, GA 30322; telephone (404) 727-

3485; fax (404) 727-4590; or email pvaleri@sph.emory.edu.

## Notice to Readers

## Epi Info 2000: A Course for Teachers of Epidemiologic Computing

CDC and Emory University's Rollins School of Public Health will co-sponsor a course, "Epi Info 2000: A Course for Teachers of Epidemiologic Computing" during March 12–15, 2001, at Emory University. The course is designed for teachers of epidemiologic computing with intermediate to advanced skills in computing.

The 4-day course covers hands-on experience with the new Windows® version of Epi Info, programming Epi Info software at the intermediate to advanced level, methods of teaching epidemiologic computing, computerized interactive exercises for teaching epidemiology, and computing. There is a tuition charge.

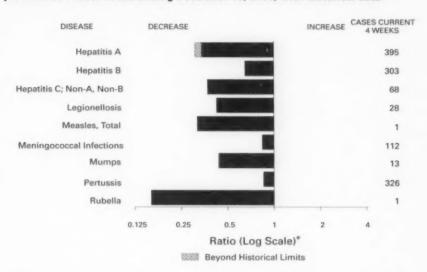
Deadline for applications is February 1. Additional information and applications are available from Emory University, Rollins School of Public Health, International Health Dept. (PIA), 1518 Clifton Road, N.E., Room 746, Atlanta, GA 30322; telephone (404) 727-3485; fax (404) 727-4590; or email pvaleri@sph.emory.edu.

# Notice to Readers

## Combined Issues of MMWR

A December 29, 2000, issue of *MMWR* will not be published. The next issue will be Volume 49, Numbers 51 and 52, dated January 5, 2001. It will include the figures and tables of notifiable diseases and deaths for the weeks ending December 23, 2000, and December 30, 2000.

FIGURE I. Selected notifiable disease reports, United States, comparison of provisional 4-week totals ending December 16, 2000, with historical data



Ratio of current 4-week total to mean of 15 4-week totals (from previous, comparable, and subsequent 4-week periods for the past 5 years). The point where the hatched area begins is based on the mean and two standard deviations of these 4-week totals.

TABLE I. Summary of provisional cases of selected notifiable diseases, United States, cumulative, week ending December 16, 2000 (50th Week)

		Cum. 2000		Cum. 2000
Anthrax			Poliomyelitis, paralytic	
Brucellosis*		61	Psittacosis*	10
Cholera		2	Q fever*	21
Cyclosporiasis	5*	3B	Rabies, human	2
Diphtheria		2	Rocky Mountain spotted fever (RMSF)	416
Ehrlichiosis:	human granulocytic (HGE)*	178	Rubella, congenital syndrome	6
	human monocytic (HME)*	98	Streptococcal disease, invasive, group A	2,619
Encephalitis:	California serogroup viral*	98 109	Streptococcal toxic-shock syndrome*	70
	eastern equine*	2	Syphilis, congenital <sup>§</sup>	70 257
	St. Louis*	3	Tetanus	26 122
	western equine®		Toxic-shock syndrome	122
Hansen diseas	se (leprosy)*	63	Trichinosis	15
Hantavirus pu	Ilmonary syndrome*1	63 30	Tularemia*	110
	emic syndrome, postdiarrheal*	165	Typhoid fever	311
HIV infection.	pediatric*1	203	Yellow fever	
Plaque		6	1.00	

: No reported cases.

\*Not notifiable in all states.

"Updated weekly from reports to the Division of Viral and Rickettsial Diseases, National Center for Infectious Diseases (NCID).

\*Updated monthly from reports to the Division of HIV/AIDS Prevention — Surveillance and Epidemiology, National Center for HIV, STD, and TB Prevention (NCHSTP). Last update November 28, 2000.

\*Updated from reports to the Division of STD Prevention, NCHSTP.

TABLE II. Provisional cases of selected notifiable diseases, United States, weeks ending December 16, 2000, and December 18, 1999 (50th Week)

	A	DS	Chie	mydia'	Countries	nasidinal-			coli O157:H	
	Cum.	Cum.	Cum.	Cum.	Cum.	Cum.	Cum.	Cum.	Cum.	LIS
Reporting Area UNITED STATES	2000° 36,091	<b>1999</b> 40,781	2000	1999	2000	1999	2000	1999	2000	Cum 1999
NEW ENGLAND	1.884		623,458	629,947	2,491	2,580	4,311	3,865	3,206	2,690
Maine	38	2,070	20,396 1,368	20,437 1,009	104 20	185 31	383	401	367	363
N.H. Vt.	31	46	1,004	944	23	19	39	35	28 35	34
Mass.	1,137	16 1,319	507 8,586	469 8,588	30	36 71	36	32	34	21
R.I.	95	96	2,409	2,251	4	6	163	176 27	168	187 26
Conn.	546	518	6,522	7,176	*	22	95	92	84	95
MID. ATLANTIC Upstate N.Y.	7,705 705	10,462 1,196	54,597	63,488	182	597	405	598	281	161
N.Y. City	3,929	5.574	23,185	25,903	128	174 256	296 12	516	72	14
N.J. Pa.	1,592	1,922	8,093	11,967	12	52	97	17 65	13 109	18 71
	1,479	1,770	23,319	25,618	31	115	N	N	87	58
E.N. CENTRAL Ohio	3,442 546	2,810	102,284	106,620	795	627	986	975	589	527
Ind.	352	317	23,724 12,648	28,285 11,595	260 58	66 41	272 133	250 104	220	219
III, Mich.	1,693	1,345	27,498	31,316	7	87	188	497	83 21	67 89
Wis.	652 199	552 134	25,179 13,235	21,439 13,985	96 374	52 381	137	124	104	83
W.N. CENTRAL	813	934	34,314				256	N	161	69
Minn.	160	177	7,129	36,573 7,238	356 131	198 75	687 236	523 167	595 211	543 186
Mo.	86 368	75 449	4,579	4,784	75	55	180	112	147	80
N. Dak.	3	6	10,975 750	12,913	33 16	26 18	103 21	46 17	97	68
S. Dak. Nebr.	7	15	1,776	1,509	15	7	56	47	20 58	18 62
Kans.	68 121	150	3,343 5,762	3,410 5,810	77	15 2	63	102	45	113
S. ATLANTIC	10,157	11,255	122,481	131,464	467	373	28 369	32	17	16
Del. Md.	199	158	2,760	2,674	6		1	332	270	188
D.C.	1,197 785	1,339 636	12,946 3,067	12,533 N	13 20	17	32	42	1	4
Va.	764	777	15,053	13,392	18	27	77	75	61	62 62
W. Va. N.C.	60 667	64 741	1,534 20,654	1,736	3	3	15	16	13	11
S.C.	755	917	9,032	17,998	28	34	90 21	74	68	52
Ga. Fla.	1,117 4,613	1,585 5,038	25,728	31,300	170	136	42	35	36	14
S. CENTRAL	1,809		31,707	30,597	209	149	90	63	76	39
(y.	186	1,788 256	47,219 7,802	44,413 7,145	49	46	147 40	141	112	104
Tenn, Ala.	771	704	14,457	13,878	11	13	61	55	32 52	35 44
Miss.	457 395	444 384	13,946 11,014	12,276	16 15	15 11	11 35	28	9	21
N.S. CENTRAL	3,708	4,159	96,162	90,166	123	90	182	9	19	4
Ark.	172	186	5,355	5,764	14	2	57	140	233 38	164 14
Okla.	649 320	814 125	17,285 8,776	15,863 7,973	10 17	24 13	9	14	53	14
Tex.	2,567	3,034	64,746	60,566	82	51	19 97	38 73	17 125	30 106
MOUNTAIN	1,322	1,605	34,774	31,714	174	100	431	328	283	242
Mont. daho	14 20	13 22	1,385 1,816	1,496 1,713	10	13	31	25		
Nyo.	9	11	769	757	23 5	8	74 21	6B 16	35 10	43
olo. I. Mex.	300 140	290 82	8,490 4,279	5,998	72	14	162	112	111	88
Ariz.	427	816	12,190	4,843 11,799	21	13	23 54	13	16	7
Jtah Vev.	137 275	141 230	2,150 3,695	2,085	28	N	52	36 35	41 70	24 48
ACIFIC	5.251	5,698		3,023	4	9	14	23	*	15
Vash.	480	336	111,231 12,606	105,072	241 N	364 N	721	427	476	398
Oreg. Calif.	171	208	5,140	5,857	21	98	222 156	164 68	200 114	180
Jaiit, Alaska	4,479	5,047	88,299 2,343	82,700	220	266	298	180	150	136
lawaii	99	93	2,843	1,817 3,086		-	30 15	14	1	1
iuam	15	18		468			N	N	U	12 U
AR.	1,245	1,180 35	3,122	U	*		7	9	Ü	Ü
mer. Samos	-	35	Ü	U	U	U	U	U	U	U
.N.M.I.	-		U	ŭ	ŭ	ŭ	Ŭ	Ü	Ü	U

N: Not notifiable.

N: Not notifiable.

U: Unavailable.

No reported cases.

C.N.M.J.: Commonwealth of Northern Mariana Islands.

Individual cases can be reported through both the National Electronic Telecommunications System for Surveillance (NETSS) and the Public Health Laboratory Information System (PHLIS).

Chiamydia refers to genital infections caused by C. trachomatis. Totals reported to the Division of STD Prevention, NCHSTP.

Updated monthly from reports to the Division of HIV/AIDS Prevention — Surveillance and Epidemiology, National Center for HIV, STD, and TB Prevention. Last update November 26, 2000.

TABLE II. (Cont'd) Provisional cases of selected notifiable diseases, United States, weeks ending December 16, 2000, and December 18, 1999 (50th Week)

	Gono	rrhea	Hepatit Non-A, I	tis C; Von-B	Legione	illosis	Listeriosis	L	yme
Reporting Area	Cum. 2000 <sup>1</sup>	Cum. 1999	Cum.	Cum.	Cum.	Cum.	Cum.	Cum.	Cun
UNITED STATES	325,596	347,165	2,838	1999 2,839	913	1999	2000	2000	199
NEW ENGLAND	5,722	6,414	16	16		1,004	647	12,874	15,12
Maine N.H.	84	78	2	2	51	78	56 2	4,313	4,46
Vt.	103	111 50	4		3	8	4	62	4 2
Mass.	2,346	2,381	4	7	5 36	14 27	3	38	2
R.I. Conn.	611 2,515	572	6	3	8	12	27	1,098 590	780
MID. ATLANTIC		3,222	~		17	14	19	2,525	3,100
Upstate N.Y.	34,181 6,862	38,472 6,558	611	123	201	242	151	6,592	8,138
N.Y. City	10,140	11,827	66	59	90	62 43	82	3,766	3,913
N.J. Pa.	5,443 11,736	7,565	510		15	21	29 21	105 1,448	1,693
E.N. CENTRAL		12,522	36	64	96	116	19	1,273	2,398
Ohio	61,766 14,321	67,164 17,443	214	888	238	265	113	325	579
nd.	5,991	6,072	12	4	111	81	58	88	44
II, Mich.	18,467 17,347	22,323	19	47	9	46 31	8	32 11	19
Vis.	5,640	14,873 6,453	182	820 16	50	64	31	-	17
V.N. CENTRAL	15,724	16,087	422		29	43	5	194	488
Minn. owa	2,784	2.741	432	301	57 7	56 13	14	421	340
Ao.	1,086 7,584	1,208 7,937	406		14	14	2	322 32	219
I. Dak.	53	79	406	287	25	18	5	44	71
S. Dak. Nebr.	270 1,320	191	-	-	2	3	2	2	1
Cans.	2,627	1,419 2,512	6	3	4 5	6	-	4	11
ATLANTIC	90,154	101,279	124	156	189	148	104	17	16
Ad.	1,671 9,003	1,615 9,635	-	-	10	19	2	974	1,288
D.C.	2,666	3,462	18	21	64	35	22	530	874
/a. V. Va.	9,891 494	9,153	3	11	33	40	8	11	6
1.C.	16,844	548 18,893	16 20	17	N	N	5	34	118
S.C.	11,071	14,479	3	22	16 6	15 11	9	46	74
la.	16,814 21,700	21,560 21,934	3 58	1 50	7	3	21	17	6
S. CENTRAL	33,995	35,316	427	331	47 37	21	37	50	33
y. enn.	3,411	3,250	36	25	20	50 22	20	47 12	99
ila.	11,469 10,959	11,120 10,857	99	117	12	22	13	28	18 57
Aiss.	8,156	10,089	8 285	188	4	4 2	4	6	20
V.S. CENTRAL	50,947	51,408	442	572	18	34	16	1	4
9.	2,920 12,870	3,159 12,672	9	28	-	1	1	45	58 5
ikla. ex.	3,968	3,905	308 10	299 16	6	11	7	4	9
	31,189	31,672	115	229	7	18	8	1 36	8 36
OUNTAIN	9,568 53	9,258 53	396	220	47	48	38	30	16
laho /yo.	89	82	3	5 8	2 5	3	-		*
olo.	52 2,688	35 2,469	303	76	2		1	3	3
. Mex.	958	943	30 16	37 34	16	13	9	11	3
rîz. tah	4,050 230	4,185 230	21	46	8	7	17		1 2
ev.	1,448	1,261	16	6	12	18	4	3	2
ACIFIC	23,539	21,767	176	232	75	83	5	4	2
/ash. reg.	2,290 766	2,047	32	21	19	21	135	127	144
alif.	19,772	857 18,129	27 115	21 190	N	N	6	15	15
laska awaii	335 376	292		130	56	60	119	101	119
uam	3/0	442 55	2		*	1	3	N	N
R.	577	313	1	2	1	*	+		
l. mer. Samoa	Ü	U	Ú	U	Ü	Ü		N	N
N.M.I.	ŭ	U	U	U	U	Ü	*	Ü	Ü

TABLE II. (Cont'd) Provisional cases of selected notifiable diseases, United States, weeks ending December 16, 2000, and December 18, 1999 (50th Week)

				00, and D	Salmonellosis*						
	Cum.	laria Cum.		s, Animal		TSS	PI	HLIS			
Reporting Area	2000	1999	Cum. 2000	Cum. 1999	Cum. 2000	Cum. 1999	Cum. 2000	Cum. 1999			
UNITED STATES	1,220	1,458	5,736	6,396	35,788	37,641	29.879	32,019			
NEW ENGLAND Maine N.H. Vt. Mass. R.I. Conn.	65 6 1 3 27 8 20	64 3 2 4 22 5 28	801 130 21 57 268 61 264	867 171 45 88 221 95 247	2,132 123 140 108 1,203 137 421	2,176 131 137 92 1,180 129 507	2,124 91 135 114 1,189 156 439	2,195 104 135 82 1,196 160 518			
MID. ATLANTIC Upstate N.Y. N.Y. City N.J. Pa.	262 81 114 36 31	415 71 246 56 42	1,112 799 U 193 120	1,267 896 U 179 192	3,938 1,190 946 820 982	5,261 1,367 1,418 1,188 1,288	4,356 1,246 886 821 1,423	5,137 1,332 1,469 1,093 1,243			
E.N. CENTRAL Ohio Ind. III. Mich. Wis.	120 22 6 46 32 14	167 18 22 76 41 10	162 52 15 22 67 6	167 36 13 10 87 21	5,036 1,530 623 1,383 869 631	5,274 1,267 526 1,566 967 948	3,375 1,350 567 176 898 384	4,590 1,008 472 1,531 948 573			
W.N. CENTRAL Minn, Iowa Mo. N. Dak, S. Dak, Nebr. Kans.	61 27 2 15 2 1 7	74 41 13 14	525 90 78 50 115 90 2 100	711 110 149 31 141 176 4 100	2,324 554 351 682 61 99 215 362	2,201 555 248 735 51 93 189 330	2,388 638 312 877 74 105 94 288	2,347 691 224 855 62 118 172 225			
S. ATLANTIC Dust. Mid. D.C. Va. W. Va, N.C. Ga.	327 5 117 16 50 4 30 2 30 67	342 1 97 18 71 4 33 15 29 74	2,325 49 401 554 112 551 155 344 159	2,078 56 389 561 108 430 133 231 170	7,989 110 803 63 983 171 1,120 740 1,477 2,522	8,542 163 831 72 1,226 168 1,295 650 1,508 2,629	5,265 137 729 U 839 148 1,072 540 1,549 251	6,340 154 871 U 1,019 150 1,282 505 1,663			
E.S. CENTRAL (y. Tenn. Alia. Miss.	47 18 12 16 1	27 7 9 7 4	199 21 102 76	252 35 93 122 2	2,356 376 656 664 660	2,188 405 571 595 617	1,656 259 755 521 121	696 1,448 291 581 479 97			
N.S. CENTRAL Ark. a. Okla. fex.	20 3 8 9	61 3 10 2 46	77 20 57	482 14 91 377	3,962 704 262 390 2,606	3,674 651 714 446 1,863	4,024 587 753 279 2,405	2,748 254 603 346 1,545			
MOUNTAIN Mont, daho Myo. Colo. N. Mex. Ariz. Jtah Nev.	52 1 4 25 9 6	44 4 3 1 18 3 7 4	247 65 9 55 21 78 10 9	215 59 5 44 1 9 81	2,797 96 128 68 692 235 841 489	2,941 82 127 68 710 361 874 527	97 51 654 182 719 451	2,555 1 97 59 693 287 804 565			
PACIFIC Nash. Oreg. Calif, Maska Hawaii	266 33 41 181	264 27 21 203 1	288	8 357 4 345 8	249 5,254 576 300 4,085 61 232	191 5,384 651 409 3,942 53 329	4,537 670 348 3,270 23 226	46 4,659 821 461 3,070 32 275			
Guam P.R. //I. Amer. Samoa C.N.M.I.	5 0 0	1 1 0 0	80 U U	70 0 0	620 U U	37 630 U U	0	00000			

N: Not notifiable. U: Unavailable. : No reported cases.

Individual cases can be reported through both the National Electronic Telecommunications System for Surveillance (NETSS) and the Public Health Laboratory information System (PHLIS).

TABLE II. (Cont'd) Provisional cases of selected notifiable diseases, United States, weeks ending December 16, 2000, and December 18, 1999 (50th Week)

			Mosis*	oo, and L	Tecember	18, 1999	(50th Week)			
		TSS		PHLIS	(Primary	yphilis & Secondary)	Tube	erculosis		
Reporting Area	Cum. 2000	Cum. 1999	Cum. 2000	Cum. 1999	Cum.	Cum.	Cum.	Cum.		
UNITED STATES	20,051	16,099	10,291	9,725	5,733	6.433	2000	1999		
NEW ENGLAND Maine N.H. Vt. Mass. R.I. Conn.	383 10 6 4 267 29 67	863 5 18 6 736 31 67	369 12 8 247 36 66	833 17 4 715 28 69	72 1 2 47 4 18	1 3 36 3	12,302 409 12 17 4 256 31	14,967 420 20 16 3 231 42		
MID. ATLANTIC Upstate N.Y. N.Y. City N.J. Pa.	2,006 760 716 337 193	1,092 281 346 270 195	1,325 211 470 384 260	722 74 233 234 181	259 15 118 48 78	16 290 19 128 67 76	2,220 284 1,195 535 206	108 2,479 307 1,273 520		
E.N. CENTRAL Ohio Ind. III. Mich. Wis. W.N. CENTRAL	3,750 409 1,506 963 642 230	3,169 410 334 1,295 507 623	1,197 309 147 103 579 59	1,754 141 113 973 453 74	1,123 69 345 350 315 44	1,200 90 427 412 230 41	1,294 263 109 628 214	379 1,588 265 132 782 310 99		
Minn. lowa Mo. N. Dak. S. Dak. Nebr. Kans.	2,369 774 522 633 51 7 142 240	1,184 233 68 697 3 18 85 80	1,884 837 316 462 49 4 84 132	793 251 58 349 2 10 68	599 133 111 27 2 6	129 9 9 93 6 12	473 165 36 192 5 16 23	515 190 54 178 6 17 16 54		
S. ATLANTIC Del. Md. D.C. Va. Vw. Va. N.C. S.C. Ga.	2,925 23 204 80 445 22 385 136 257 1,373	2,363 15 160 51 130 8 206 119 230 1,444	1,110 23 115 U 331 17 265 87 167 105	525 10 98 U 64 5 92 63 83 150	1,918 8 289 48 126 2 469 212 370 394	2,040 8 337 46 150 5 449 248 435 363	2,598 14 239 36 258 31 390 128 555	3,070 26 258 52 268 37 482 222 568		
E.S. CENTRAL Ky. Tenn. Ala. Miss.	1,134 495 340 98 201	1,182 232 659 117 174	525 112 357 49 7	677 147 457 62 11	851 82 513 122 134	1,114 101 631 202 180	947 852 114 305 296	1,157 999 176 346 295		
W.S. CENTRAL Ark. .a. Okla. Fex.	2,893 203 138 125 2,427	2,602 74 217 514 1,797	2,606 52 191 43 2,320	1,162 26 134 160 842	802 94 204 126 378	1,022 86 300 182 455	137 1,022 159 74 130 659	182 1,766 166 245 176 1,179		
MOUNTAIN Mont. daho Wyo. Colo. N. Mex. Ariz. Jtah	1,286 8 45 5 266 168 594 80 120	1,118 10 27 3 198 145 573 64	732 25 3 196 99 329 80	756 12 1 156 105 410 66	225 1 1 11 21 185 1	230 1 1 1 4 11 206 2	471 17 13 4 70 36 214	516 13 15 3 75 61 219		
PACIFIC Nash. Oreg. Calif. Alaska tawaii	3,306 447 163 2,648 8 36	98 2,526 121 94 2,274 4 33	543 405 105 3 30	6 2,503 111 86 2,268 4 34	5 424 67 6 349	350 66 7 274 1 3	71 2,963 236 25 2,481 96 125	3,614 239 104 3,034 59 178		
Guam P.R. /I. Amer. Samoa C.N.M.I.	35	19 138 U U	ככככט	0000	159 U U	144 U U	119	70 178 U		

N: Not notifiable.

U: Unavailable.

No reported cases.

Individual cases can be reported through both the National Electronic Telecommunications System for Surveillance (NETSS) and the Public Health Laboratory Information System (PHLIS).

TABLE III. Provisional cases of selected notifiable diseases preventable by vaccination, United States, weeks ending December 16, 2000, and December 18, 1999 (50th Week)

	H. influ	ienzae,	H	epatitis (V	iral), By Ty	pe			Measl	es (Rubec	ofa)	
		sive	A	A			Indiger	ious	Imported*		Total	
Reporting Area	Cum. 2000'	Cum. 1999	Cum. 2000	Cum. 1999	Cum. 2000	Cum. 1999	2000	Cum. 2000	2000	Cum. 2000	Cum. 2000	Cum. 1999
NITED STATES	1,168	1,170	11,963	15,786	6,417	6,662	-	61	-	18	79	94
EW ENGLAND	102	97	352	340	97	139	-	3		4	7	11
laine	2	8	21	14	5	1	-		-	*	-	
I.H.	12	18	10	17	18	16	-	2	~	1	3	1
t. Aass.	10 40	5 39	10	19 138	6 18	4		1	-	3	3	8
I.I.	4	6	25	26	22	33		-	-		-	
onn.	34	21	156	126	28	41	U	-	U	*		2
AID. ATLANTIC	183	198	1,071	1,140	836	865	-	14	*	5	19	5
Ipstate N.Y.	98	81	220	264	137	176	+	9	-		9	2
I.Y. City	42 33	57 53	375 100	385 145	428 57	275 135	-	5	-	4	9	3
a.	10	7	376	346	214	279	-	-	-	1	1	-
N. CENTRAL	150	194	1,497	2,867	693	687		9			9	4
Ohio	53	59	264	640	101	90		2	-	-	2	
nd.	30	25	119	102	46	42	*		-	-	-	2
II. Aich.	54 10	83	623 478	816 1,235	110 435	52 473		4 3	-	-	3	1
Mich. Vis.	3	7	13	74	435	30		3		-	3	
V.N. CENTRAL	74	75	699	1,025	526	347		3		2	5	1
Minn.	43	47	184	96	41	52	-	3		1	1	1
owa	1	2	65	141	32	41	-	2		-	2	
Mo.	18	11	301	667	381	215	-	-	-	~	-	-
N. Dak. S. Dak.	4	1 2	4 3	3 9	2 2	2		-				
Nebr.	3	4	34	49	44	20				*	-	
Cans.	4	8	108	61	24	16	-	1		1	2	
S. ATLANTIC	292	251	1,505	1,815	1,299	1,097	-	4		-	4	20
Del. Vid.	75	68	221	299	120	145	-	-	-	*	2	
D.C.		5	35	59	34	25	-	-	-	-	-	
Va.	39	22	154	175	162	97	-	2	-	-	2	18
W. Va. N.C.	9 23	36	55 149	40 160	21 246	23	-	-	*	*	-	-
S.C.	15	6	86	46	23	63		-	- 2			
Ga.	70	GB	289	452	222	166		-	-	9	-	-
Fla.	61	39	516	582	471	365	-	2		*	2	2
E.S. CENTRAL	51	67	383	394	456	465		-	-	*		2
Ky.	12	38	48 140	66 147	75	46 207	-	-	7	*	-	2
Tenn. Ala.	26 12	18	57	60	218 56	86		-		-	- 1	
Miss.	1	3	138	121	107	126	-	-	-	-		
W.S. CENTRAL	58	61	2,224	2,962	706	1,104				2.		12
Ark.	2	2	112	74	78	84	-	-	-	-	-	5
La.	11	15	60	212	93	171	-	-	-	-	+	
Okla. Tex.	43	40	256 1,796	489 2,187	154 381	145 704		-		-		7
MOUNTAIN	117	105	987	1,213	561	551		12		1	13	2
Mont.	1	3	7	1,213	7	17		12			13	4
ldaho	4	1	42	45	8	29	-	-	-	-	*	
Wyo.	1 21	1 14	45	9	38 110	14 98	-	-		-	-	
Colo. N. Mex.	24	19	207 70	215 51	113	174		2	~	1	3	
Ariz.	49	54	474	670	210	131			-		-	1
Utah	11	9	64	63	27	34		3	-	-	3	
Nev.	6	4	78	143	48	54	-	7		-	7	1
PACIFIC	141	122	3,245	4,030	1,243	1,407		16	*	6	22	37
Wash. Oreg.	8 29	8 41	279 177	387 244	113 120	76 114	-	2	-	1	3	12
Calif.	33	53	2,765	3,362	989	1,186	-	13	-	2	15	17
Alaska	45	9	11	13	10	16		1	-	100	1	
Hawaii	26	11	13	24	11	15		-	-	3	3	3
Guam P.R.	4	2	230	348	259	248	U	-	U		7	
V.I.	ű	ű	U	U	739 U	240 U	U	U	U	Ú	ú	ı
Amer. Samoa	U	U	U	U	U	U	U	U	U	U	U	L
C.N.M.I.	U	U	U	U	U	U	U	U	U	U	U	- 1

N: Not notifiable.

- : No reported cases.

- : No rep

TABLE III. (Cont'd) Provisional cases of selected notifiable diseases preventable by vaccination, United States, weeks ending December 16, 2000, and December 18, 1999 (50th Week)

		jococcal ease		Mumps			Pertussis			Rubella	
Reporting Area	Cum. 2000	Curn. 1999	2000	Cum. 2000	Cum. 1999	2000	Cum. 2000	Cum. 1999	2000	Cum. 2000	Cum. 1999
JNITED STATES	1,966	2,243	4	308	364	68	6,368	6,400	2000	151	249
NEW ENGLAND	122	108	-	4	9	6	1,566	861		13	7
Aaine I.H.	8 12	5 12			2	-	45 125	96		2	
t. Aass.	3 72	5 63	-	1	1 4	4	244 1,085	86 609	-	9	7
1.1.	9	7		1	2	2	22	38		1	-
conn.	18	16	U	2	*	U	45	32	U	1	
AID. ATLANTIC Spstate N.Y.	189 66	222 71		24 11	44 12	2 2	616 313	1,020 764	7	9 2	35 21
N.Y. City N.J.	37 44	56 50	-	4 3	12	-	51 42	60 27	-	7	7 4
a.	43	46	-	6	18	-	210	169		-	3
.N. CENTRAL	350	396	-	30	51	4	725	654	-	1	2
Ohio nd.	94 48	131 64	-	7	20 5	3	321 119	291 82	-	-	1
II. Mich.	78 104	104 61	-	6	13	1	79 124	97 70	7	1	1
Nis.	26	36	-	-	4	-	82	114	+	-	-
V.N. CENTRAL	149	222 48	1	19	14	16	585	484		3	130
Vlinn. owa	21 34	38	-	7	8	14	365 55	226 95	-	1	30
Mo. N. Dak.	68	89	-	4	1	-	79	73 18	-	1	2
S. Dak.	6	11	-	:	-	-	7	7	-	3	-
Nebr. Cans.	10	11 21	1	4	3	2	32 40	9 56	-	1	92
S. ATLANTIC	308	380	2	48	49	11	503	430		96	37
Del. Md.	26	10 53	-	10	6	7	122	119		1	1
D.C. Va.	42	4 56	1	11	10	1	112	51	-	-	
N. Va.	12	8	-	7		-	1	4			
N.C. S.C.	36 26	47	-	11	8	2	110 41	101		82 10	36
Ga. Fla.	118	61 98	1	7	14	i	40 66	40 90	-	2	-
E.S. CENTRAL	130	156	1	8	15	1	106	111		5	2
Ky. Tenn.	26 56	33 64		1 2	-	1	54 32	42 45	- 1	1	
Ala. Miss.	36 13	36 23	1	3 2	11		19	21	-	3	2
W.S. CENTRAL	131	207		31	46	1	338	215	-	6	15
Ark.	14	35		5			36	25	-	-	5
La. Okla.	35 28	66 35	2	4	11	1	12 41	9		1	1
Tex.	54	71	-	22	32	-	249	140	-	5	9
MOUNTAIN Mont.	165 6	137	-	26	26	13	796 36	774	-	2	16
ldaho	7	12	-	1	3	-	64	145		-	,
Wyo. Colo.	34	5 36	1	4 2	6	-	6 457	287	-	1	1
N. Mex. Ariz.	12 91	15 41	-	1 4	N 8	1	89 99	149 118	-	1	13
Utah	8	16	-	7	4		31	58	-	-	1
Nev.	4	8	-	6	5	-	15	13	-		1
PACIFIC Wash.	422 64	415 65		118	110	14 12	1,133 416	1,851 645		17	5
Oreg. Calif.	75 266	76 259	N	N 86	N 89	2	113 550	1,091	-	10	
Alaska	9	7	-	7	3	*	22.	5	-	-	
Hawaii	8	8		14	16	-	32	50	11	-	
Guam P.R.	9	13	U	0	-	U	7	31	U	-	
V.I. Amer. Samoa	U	Ü	U	U	U	U	U	U	U	U	U
C.N.M.I.	ŭ	ŭ	ŭ	ŭ	ŭ	ŭ	ŭ	ŭ	ŭ	ŭ	ŭ

N: Not notifiable. U: Unavailable. -: No reported cases.

# TABLE IV. Deaths in 122 U.S. cities,\* week ending December 16, 2000 (50th Week)

	-	All Cau	ses, By	Age (Y	ears)		PBd			All Cau	ses, By	Age (Y	ears)		P&I'
Reporting Area	All Ages	≥65	45-64	25-44	1-24	<1	Total	Reporting Area	All Ages	≥65	45-64	25-44	1-24	<1	Total
NEW ENGLAND Boston, Mass. Aridgeport, Conn. Ambridge, Mass. All River, Mass. Hartford, Conn. Lowell, Mass. Lynn, Mass. New Bedford, Ma New Haven, Conn Providence, R.I. Somerville, Mass. Springfield, Mass.	14 30 50 25 13 ss. 26 U 2	3677 689 386 100 211 322 200 100 244 311 U.	28 4 2 5 10 4 2 2 2 11 U	53 32 2 2 4 5 1	16 11 2 1 1 U	15 10 1 	80 18 3 3 - 2 7 3 3 5 U	S. ATLANTIC Atlanta, Ga. Baltimore, Md. Charlotte, N.C. Jacksonville, Fla Miami, Fla. Norfolk, Va. Richmond, Va. Savannah, Ga. St. Petersburg, Tampa, Fla. Washington, D. Wilmington, De	92 54 56 40 Fla. 82 C. 100 I. 25	39 31 64 143 57 18	266 34 60 14 37 21 10 13 6 8 30 26 7	102 18 22 7 15 7 5 3 1 9 6	35 4 6 4 3 3 1 1 8 5	17 1 4 3 1 1 2 2 3	79 4 17 9 7 12 1 5 4 4 15
Waterbury, Conn. Worcester, Mass. MID. ATLANTIC Albany, N.Y. Allentown, Pa. Buffalo, N.Y. Camden, N.J. Eirabeth, N.J. Erie, Pa. §	2,250 37 15 99 29 23 50	1,588 30 10 70 22 16 40	13 462 5 1 2 16 4 5 7	3 136 1 5 2 1 2	36 1 2 1	24	11 134 3 1 12 4	E.S. CENTRAL Birmingham, Al Chattanooga, Te Knoxville, Tenn. Lexington, Ky. Memphis, Tenn Mobile, Ala. Montgomery, A Nashville, Tenn.	enn. 80 114 66 . 185 96 la. 60	118 64 85 46 134 72 36	152 31 12 20 11 16 12 11 39	64 11 3 9 8 7 8 5	32 2 2 12 5 7 2	27 2 1 2 16 1 1 4	68 16 4 10 4 11 5 5
Jersey City, N. J. New York City, N. Newark, N. J. Paterson, N. J. Philadelphia, Pa. Pittsburgh, Pa. S Reading, Pa. Rochester, N. Y. Scranton, Pa. § Syracuse, N. Y. Trenton, N. J. Utica, N. Y. Yonkers, N. Y.	66 32 282 46 34 117	91 21 6	259 18 9 63 63 5 4 14 7 7 3 2 1 22 3	3 74 13 3 13 3 2 9 1	1 13 4 1 4 2 1 1	1 10 4 3 2 1	55 5 3 20 1 5 7 3 2 4 3 3 U	W.S. CENTRAL Austin, Tex. Baton Rouge, Li Corpus Christi, Dallas, Tex. El Paso, Tex. Houston, Tex. Little Rock, Ark. New Orleans, Li San Antonio, Te Shreveport, La. Tulsa, Okla.	Tex. 54 183 65 103 1 44 1. 7	68 25 42 122 46 70 1 U 25 42 174 12	206 16 7 7 28 11 23 U 17 9 56 2 31	89 11 4 1 25 7 5 U 2 11 15	31 3 1 2 3 1 1 U 3 4 9	28 4 2 5 3 4 U 2 3 4 1	71 10 4 14 3 2 2 U 2 14 10 4 8
E.N. CENTRAL Akron, Ohio Canton, Ohio Chicago, III. Cincinnati, Ohio Cleveland, Ohio Columbus, Ohio Dayton, Ohio Detroit, Mich. Evansville, Ind. Fort Wayne, Ind.	2,113 49 42 354 170 137 163 144 236 25	3 3 24 11 10 10 11 11 13 2	8 6 2 7 1 65 8 44 4 20 7 23 2 23 1 63 5 4	2 28 6 7 11 6 23	51 1 1 11 1 1 5 2 14	9 1 5 7 1 5	143 3 6 25 12 5 9 12 9 7 6	MOUNTAIN Albuquerque, N Boise, Idaho Colo. Springs, ( Denver, Colo. Las Vegas, Nev. Ogden, Utah Phoenix, Ariz., Pueblo, Colo. Salt Lake City, L Tucson, Ariz.	2010. 8 110 243 201	U U 26 1 46 1 46 1 54 3 154 3 20 1 133 U U	U 4 17 23 65 4 33 U	13 5 16 2 13 U 9 8	32 3 2 7 9 U 5 6	28 U 1 2 7 1 11 U 6	6E U
Gary, Ind. Grand Rapids, Mi Indianapolis, Ind Lansing, Mich. Milwaukee, Wis. Peoria, III. Rockford, III. South Bend, Ind. Toledo, Ohio Youngstown, Oh	205 225 115 50 61 100	1 13 1 2 1 8 1 3 1 4	0 2 4 6 8 43 1 3 1 21 5 10 9 9 U U 8 13	3 6 15 3 8 4 2 U	1 2 5 1 1 U 2 2	3 4 1 4 1 1 U	2 5 18 1 12 2 2 U 5 2	PACIFIC Berkeley, Calif. Fresno, Calif. Glendale, Calif. Honolulu, Haw. Long Beach, Ca Los Angeles, Ca Pasadena, Calif Portland, Oreg. Sacramento, Ca	lif. 8 slif. ( 14	U U U U U U U U U U U U U U U U U U U	U U 7 16 U 3 28	41 U U 2 4 U 1 3 U	17 U U U	16 0 0 0 1 1 1 0 1 1 0	91
W.N. CENTRAL Des Moines, low Duluth, Minn. Kansas City, Kan. Kansas City, Mo. Lincoln, Nebr. Minneapolis, Mi. Omaha, Nebr. St. Louis, Mo. St. Paul, Minn.	50 5. 25 103 36	7 0 4 5 1 8 6 8 3 7 9 7 7	1 135 3 16 1 6 6 7 4 26 2 3 3 21 7 13 16 17	4 3 2 5 2 7 4 22	19 2 4 5 2 1 2	18 1 4 1 1 1 6 2 2	59 10 3 2 13 2 6 10	San Diego, Cali San Francisco, San Jose, Calif Santa Cruz, Cal Seattle, Wash. Spokane, Wash Tacoma, Wash.	f. 18 Calif. 14 . I if. 3	2 135 4 97 J U 4 22 4 98	31 28 U 11 20 8 U	12 9 U 1 4 5 U	2 4 U 4 1 U 269	1 5 U 7 U 217	2 11 1

U: Unavailable. ∴No reported cases.
\*Mortality data in this table are voluntarily reported from 122 cities in the United States, most of which have populations of ≥100,000.
A death is reported by the place of its occurrence and by the week that the death certificate was filed. Fetal deaths are not included.
\*Peneumonia and influenza.
\*Because of changes in reporting methods in this Pennsylvania city, these numbers are partial counts for the current week. Complete counts will be available in 4 to 6 weeks.
\*Total includes unknown ages.

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